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BK channels of five different subunit combinations underlie the de novo KCNMA1 G375R channelopathy

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The molecular basis of a severe developmental and neurological disorder associated with a de novo G375R variant of the tetrameric BK channel is unknown. Here, we address this question by recording from single BK channels expressed to mimic a G375R mutation heterozygous with a WT allele. Five different types of functional BK channels were expressed: 3% were consistent with WT, 12% with homotetrameric mutant, and 85% with three different types of hybrid (heterotetrameric) channels assembled from both mutant and WT subunits. All channel types except WT showed a marked gain-of-function in voltage activation and a smaller decrease-of-function in single-channel conductance, with both changes in function becoming more pronounced as the number of mutant subunits per tetrameric channel increased. The net cellular response from the five different types of channels comprising the molecular phenotype was a shift of −120 mV in the voltage required to activate half of the maximal current through BK channels, giving a net gain-of-function. The WT and homotetrameric mutant channels in the molecular phenotype were consistent with genetic codominance as each displayed properties of a channel arising from only one of the two alleles. The three types of hybrid channels in the molecular phenotype were consistent with partial dominance as their properties were intermediate between those of mutant and WT channels. A model in which BK channels randomly assemble from mutant and WT subunits, with each subunit contributing increments of activation and conductance, approximated the molecular phenotype of the heterozygous G375R mutation.

Introduction

The BK channel (Slo1, KCa1.1) is a large conductance K⁺ selective channel that is synergistically activated by Ca²⁺ and voltage (Barrett et al., 1982; Latorre et al., 1982, 2017; McCobb et al., 1995; Rothenberg and Magleby, 2000; Horrigan and Aldrich, 2002; Xia et al., 2002; Salkoff et al., 2006; Horrigan, 2012; Geng and Magleby, 2015; Pantazis and Olcese, 2016; Hite et al., 2017; Tao et al., 2017; Zhou et al., 2017; Tao and MacKinnon, 2019; Geng et al., 2020). BK channels are homotetrameric proteins comprised of four large pore-forming (α) subunits >1,200 amino acids, encoded by the KCNMA1 gene (Fig. S1). BK channels are widely expressed in many cell types where they modulate smooth muscle contraction (Brenner et al., 2000), transmitter release (Robitaille et al., 1993), circadian rhythms (Harvey et al., 2020), repetitive firing (Gu et al., 2007; Park et al., 2022), and cellular excitability (Montgomery and Meredith, 2012). Mutations in the KCNMA1 gene that encode the (α) subunit of BK channels are associated with a wide range of diseases, including epilepsy, dyskinesia, autism, multiple congenital abnormalities, developmental delay, intellectual disability, axial hypotonia, ataxia, cerebral and cerebellar atrophy, bone thickening, and tortuosity of arteries (Wang et al., 2009; Yang et al., 2010; Bailey et al., 2019; Liang et al., 2019; Cui, 2021; Miller et al., 2021).

Studies of the pathogenic properties of BK channels associated with diseases have often been incomplete, focusing on the homotetrameric mutant channels. Yet, for a mutation heterozygous with the wild-type (WT) allele, mutant and WT subunits have the potential to assemble into five different stoichiometries for tetrameric channels with likely differences in functional properties (Fig. 1; MacKinnon, 1991; Blaine and Ribera, 1998; Niu and Magleby, 2002; Bergendahl et al., 2019; Backwell and Marsh, 2022).

Here, we show that this is the case for a de novo G375R mutation in the (α) subunit of BK channels associated with the Liang-Wang Syndrome (Liang et al., 2019). Three unrelated children who carried this mutation had a syndromic neurodevelopmental disorder associated with severe developmental delay and polymalformation syndrome (Liang et al., 2019). The G375R mutation, located on the back side of the S6 pore-lining helix of the α subunit (Fig. S1), replaces the hydrogen side chain of glycine with a large arginine side chain that might be
expected to distort the subunit structure and gating (Chen et al., 2014) as well as add a positive charge that may alter conductance.

To assess the functional effects of a G375R mutation, we recorded currents from whole cells and excised macropatches of the membrane after injecting a 1:1 mixture of G375R mutant and WT cRNA encoding mutant and WT subunits into *Xenopus laevis* oocytes to mimic a de novo mutation heterozygous with a WT allele. When compared with WT, the currents from the 1:1 injection were left-shifted more than $-120$ mV. This large negative shift in activation indicated a pronounced gain-of-function (GOF) mutation at the cellular level causing the channels to open inappropriately at negative membrane voltages. These observations provide a possible explanation for the severity of the disease associated with the heterozygous G375R mutation.

To investigate the molecular basis underlying the cellular response, detailed single-channel recording (Hamill et al., 1981) following the 1:1 injection suggested that five different types of functional BK channels were expressed: 3% were consistent with WT, 12% with homotetrameric mutant, and 85% with three different types of hybrid (heterotetrameric) channels. All channel types except WT showed a marked GOF in voltage activation and a smaller decrease-of-function (DOF) in single-channel conductance, with both becoming more pronounced as the number of mutant subunits per channel increased. Codominance was observed for the two homotetrameric channels, with homotetrameric WT channels active at the most positive voltage range of channel activity and homotetrameric mutant channels active at the most negative. Partial dominance was observed for the three types of hybrid channels, which were activated at voltages intermediate between those of WT and homotetrameric mutant channels. A model in which BK channels were randomly assembled from mutant and WT subunits, with each subunit contributing increments of activation and conductance, could approximate the molecular phenotype of the heterozygous G375R BK mutation. The possibility that a channelopathy patient with a heterozygous BK channel mutation synthesizes five different types of BK channels in their neurons and other cells, four with aberrant properties, presents a daunting challenge for treatment.

### Materials and methods

#### Constructs

Experiments were performed using the human large conductance calcium-activated potassium (BK) channel (hSlo1) KCNMA1 transcript: GenBank accession no. U23767.1 (McCobb et al., 1995) for WT, and a G375R mutation (VFFILGGLAMF to VFFILRGLAMF) was constructed to match the G375R de novo variant described by Liang et al. (2019). G375R is at the same position as G310 in Tao and MacKinnon (2019), which they called the gating hinge residue. The numbering differs because...
Tao and MacKinnon (2019) used an alternative transcript missing the first 65 amino acids compared to U23767.1. Both our hSlo1 WT cRNA plasmid and WT mammalian transfection plasmid contain the same hSlo1 insert and are described in McCobb et al. (1995). Overlap extension PCR cloning was used to generate the G375R mutation, which was verified by sequencing. The new construct was linearized downstream of the end of coding and transcribed with T3 using Invitrogen’s T3 mMessage mMachine kit to make cRNA for injection into Xenopus oocytes. To mimic the effects of a heterozygous mutation, G375R mutant and WT cRNA were mixed in a 1:1 molar ratio before injection into oocytes. For expression in HEK293 cells, two mammalian transfection plasmids were created. DNA fragments containing the complete channel cDNAs of either the WT or G375R mutant and two unique restriction sites flanking it were amplified by PCR and ligated into mammalian transfection plasmids, which were verified by sequencing.

Expression of BK channels in oocytes for whole-cell recording
Defolliculated Xenopus oocytes were injected with 0.5–150 ng of cRNA using a Nanoject II (Drummond Scientific) and incubated at 18°C for 2–5 d before recording. Incubation was in ND96 complete medium consisting of (in mM) 96 NaCl, 2 KCl, 1.8 CaCl2, 5 MgCl2, and 5 HEPES, adjusted to pH 7.5, supplemented with 2.5 mM sodium pyruvate and 100 µg/ml each penicillin and streptomycin. The two-microelectrode whole-cell voltage-clamp recordings from oocytes were obtained in ND96 medium with 1 mM added 4,4’-diisothiocyanato stilbene-2,2’-disulfonic acid disodium salt hydrate (DIDS) to block the endogenous chloride conductance. Currents were obtained with an Oocyte Clamp OC-725C amplifier (Warner Instrument Corp.). Recordings were low-pass filtered at 1 kHz and digitized at 10 kHz. Electrodes were made with borosilicate glass capillaries (World Precision Instruments) pulled with a Sutter Instrument Co. P-87 pipette puller and filled with 3 M KCl.

Macropatch and single-channel recordings from BK channels expressed in Xenopus oocytes
Oocytes were injected with 0.1–18 ng of cRNA and incubated at 18°C for 2–5 d in Barth’s Solution (in mM): 88 NaCl, 1 KCl, 2.4 NaHCO3, 0.33 Ca(NO3)2, 0.41 CaCl2, 0.82 MgSO4, 15 mM HEPES, pH 7.6, plus 12 µM tetracycline. Macroc- and single-channel currents were recorded from inside-out patches of the membrane (Hamill et al., 1981) excised from oocytes at room temperature (21–24°C). pClamp 9.0 software (Molecular Devices) was used to drive an Axopatch 200B amplifier to collect the currents. For macropatch current recordings, borosilicate pipettes with 0.5–2 MΩ resistance were used. The macrocurrennts were filtered at 10 kHz and sampled at 100 kHz. A minus P/4 protocol was used to remove capacitive transients and leak currents. For single-channel recordings, borosilicate pipettes with 8–12 MΩ resistance were used. The single-channel currents were filtered at 5 kHz and sampled at 200 kHz. The pipette (external) solution contained (in mM) 160 KCl, 2 MgCl2, and 5 TES buffer, pH 7.0. The internal membrane surface of the excised patches was perfused by two different solutions. The designated 0 Ca2+ solution had a free Ca2+ <0.01 µM and contained (in mM) 160 KCl, 1 EGTA, 1 HEDTA, and 10 HEPES, pH 7.0. The solution with 300 µM internal Ca2+ contained (in mM) 160 KCl, 0.3 CaCl2, and 10 HEPES, pH 7.0. Procedures to obtain oocytes from Xenopus laevis were approved by the University of Miami Animal Care and Use Committee. Macropatch and single-channel recordings were analyzed with Clampfit 10.7 software (Molecular Devices) and SigmaPlot 12.

For injection of only G375R cRNA into Xenopus oocytes, we found that it was difficult to get giga-ohm seals of sufficient quality for macropatch recordings using the lower resistance electrodes required for such recordings. Hence, macropatch currents are not presented for injection of only G375R cRNA into oocytes. However, it was still possible to obtain high-quality giga-ohm seals for single-channel recording using the higher resistance pipettes required for such recordings. We also found that the viability of the oocytes was greatly reduced following injection of only G375R cRNA, with the oocytes often starting to die by the second or third day after injection, rather than after a week or more, perhaps because a large fraction of the G375R homotrameric mutant channels would be expected to be open at resting membrane potentials, as will be shown in later sections. These viability problems were not observed for injection of 1:1 mixtures of G375R mutant and WT cRNA, which gave less pronounced negative shifts in activation, or for injections of only WT cRNA.

The open probability (Po) of each single channel analyzed in detail was determined for a range of voltages with Clampfit 10.7 using 50% threshold analysis to measure open and closed interval durations. Po at each voltage was calculated by dividing the total open time by the sum of the open and closed times. The duration of recordings to estimate Po ranged from ~5 s to ~3 min, with the time increasing as the Po decreased. For macropatch current recordings, relative conductance was determined from macroscopic tail current amplitudes using the voltage protocols indicated in the figure legends. G/Gmax vs. V (G-V) plots for macrocurrent recordings and Po vs. V (Po-V) plots for single channels were fitted with a Boltzmann function to estimate voltage for half activation (V1/2), the voltage required for half-maximal activation, and b, a measure of voltage sensitivity, using

\[
G/G_{\text{max}} = \frac{1}{1 + \exp[(V_i - V)/b]},
\]

where G/Gmax is the ratio of conductance to maximum conductance and b is the slope factor which gives a measure of voltage sensitivity, where b indicates the change in millivolts required to increase G/Gmax (or Po for single channel recording) e-fold at very low G/Gmax (or Po). Note that an increase in b indicates a decrease in slope and voltage sensitivity.

Single-channel conductance was determined at 100 mV by setting horizontal cursor lines by eye to the open and closed current levels of single-channel recordings.

HEK293 cell culture
As described previously (Santi et al., 2006), human embryonic kidney HEK293 cells (ATCC CRL-1573; ATCC) were cultured in Gibco Dulbecco’s modified Eagle’s medium (DMEM) with 10% fetal bovine serum, 100 units ml⁻¹ penicillin, and 100 µg ml⁻¹.
streptomycin (Thermo Fisher Scientific) and incubated at 37°C with 5% CO2. These cells were grown and passaged twice a week in T25 flasks (MidSci).

Expression of BK channels in HEK293 cells
HEK293 cells were plated at a density of ~400,000 cells per 40-mm petri dish (#93040; TPP catalog) a day before transfection. For transfection, plasmid DNA containing cDNA encoding for BK WT and/or BK G375R (mutant) subunits was added to cell layers that were 70–90% confluent using Lipofectamine 2000 transfection reagent (catalog #11668-027; Thermo Fisher Scientific) following manufacturer’s instructions. 2 μg cDNA of WT, mutant, or a mix of WT and mutant (1:1 ratio) was transfected per 40 mm dish. As a marker for transfection, cells were co-transfected with pmaxGFP, a CMV plasmid expressing green fluorescent protein (Amaxa Biosystems) at 0.2 μg per dish. After transfection, the cells were incubated at 37°C with 5% CO2 for 2–4 d until recording. The recordings were then performed at room temperature (22°C). Fluorescent cells were used to study the function of the ion channels that the HEK293 cells were transfected with; non-fluorescing cells were used as a non-transfected control (Han et al., 2004).

Whole-cell recording from HEK293 cells
Whole-cell recordings were obtained from HEK293 cells using an Axopatch 200B amplifier (Molecular Devices). Recordings were filtered at 5 kHz with the amplifier internal filter and digitized at 50 kHz using a Digidata 1550B digitizer (Molecular Devices). The recording pipette was filled with (in mM) 140 KMES, 5 EGTA, and 10 HEPES, pH 7.4 with KOH. Bath solutions contained (in mM) 135 NaMES, 5 KMES, 1 MgCl2, and 10 HEPES, pH 7.4 with NaOH.

Calculating the percentages of the five different stoichiometries of BK channels that can be expressed when mimicking a heterozygous mutation
As shown in Fig. 1, a mutation heterozygous with WT can potentially produce five different types of channels defined by their stoichiometry. The binomial equation was used to calculate the expected percentages of different combinations of mutant and WT subunits for tetrameric BK channels (Blaine and Ribera, 1998; MacKinnon, 1991). For the calculations in Fig. 1, this approach assumes that WT and mutant alleles produce equal numbers of mutant and WT subunits that assemble randomly into tetrameric channels of different stoichiometries, labeled assembled channels in Fig. 1, and that each assembled channel has the same probability of being expressed, independent of subunit composition, such that

\[ F_i = \frac{\binom{n}{i}}{\binom{n}{i} + \binom{n}{i}} F_{WT}^{(n-i)} F_{MT}^{i} \]  

(2)

where \( F_i \) is the decimal fraction of channels with \( i \) mutant subunits, \( F_{WT} \) is the fraction of WT cRNA injected, and \( F_{MT} \) is the fraction of mutant cRNA injected, where \( F_{WT} + F_{MT} = 1 \). For BK channels, \( i \) ranges from 0 to 4, giving five channel types based on subunit composition. Setting both \( F_{WT} \) and \( F_{MT} \) to 0.5 gives the fractions of the various channel types for a heterozygous mutation. \( F_i \) multiplied by 100 gives the percentage of expressed channels of each type.

Simulating discrete probability distributions of the number of BK channels of each stoichiometry expected for equal production and random assembly of mutant and WT subunits
Simulated discrete probability distributions were used to determine if the experimentally observed number of channels of each stoichiometry in a group of 33 assembled channels expressed following 1:1 injection of mutant and WT cRNA differed significantly from the numbers expected assuming equal production and a random assembly of mutant and WT subunits. The discrete probability distributions for each of the five types of channels based on subunit stoichiometry were simulated as follows. The first step was to generate a group of 33 channels by assuming a random assembly of mutant and WT subunits drawn from equal numbers of mutant and WT subunits (with replacement) for each of the 33 channels in the group. To assemble each of the 33 channels, four random numbers between 0 and 1 were drawn. Each random number <0.5 indicated a WT subunit and each random number ≥0.5 indicated a mutant subunit. The subunit composition of each simulated channel then indicated its channel type from the stoichiometry. The number of channels of each type in each group of 33 channels was tabulated and binned into five separate frequency histograms, one for each channel type. To give an example for one group of 33 channels, the simulation may have assembled 1 channel with four WT subunits, 10 channels with one mutant and three WT subunits, 11 channels with two mutant and two WT subunits, 7 channels with three mutant and one WT subunits, and 4 channels with four mutant subunits. The frequency histogram indicating the number of WT channels in the groups of 33 channels would then have one count added to the specific bin indicating 1 WT channel; the frequency histogram indicating the number of channels with one mutant and three WT subunits found in groups of 33 channels would then have one count added to the bin indicating 10 channels of that type; the frequency histogram indicating the number of channels with two mutant and two WT subunits found in the group of 33 channels would then have one count added to the bin indicating 11 channels of that type, with similar binning for the 2 remaining channel types. This process was repeated for 106 groups of 33 channels, where one count was added to each of the five frequency histograms for each group of 33 channels. If no channels of a given type were observed in a group of 33 channels, then the count was added to bin 0 to indicate that 0 channels of that type were observed in the group of 33 channels. Each frequency histogram for each channel type then contained 106 total counts. Dividing the number of counts in each bin in the frequency histograms by 106 counts in that frequency histogram gave the discrete probabilities of observing 0, 1, 2, 3...33 channels of the indicated stoichiometry in a group of 33 channels. The discrete probabilities summed to 1.0 for each distribution. The five discrete probability distributions, one for each of the five types of assembled channels, are presented in Fig. S4.

Software
The increments of single-channel conductance and \( V_b \) added by each G375R mutant and WT subunit for Fig. 5 C and Fig. 6 D were optimized by minimizing the sum of the squared differences...
similarly in neurons, the flux of K+ through opened BK channels would act to drive the membrane potential toward the equilibrium potential for K+, opposing depolarization of the cell and facilitating repolarization, both of which could interfere with normal neuronal function. Thus, at the level of whole-cell recording, the heterozygous G375R mutation gave a pronounced GOF phenotype because much less depolarization was required to activate BK currents. Some of the disease phenotypes

between the experimental values and those calculated with the equations in the legends of Fig. 5 C and Fig. 6 D by running Solver in Excel with solving method GRG nonlinear. The computer program for simulating the discrete probability distributions for observing a given number of channels of a given type in a group of 33 channels assuming random assembly of mutant and WT subunits, Binomdouble5-33.bas, was written in QB64 and is available from K.L. Magleby upon request.

Statistics
Error bars in the figures and error estimates in the text are SEM. Significance was determined with the two-tailed t test unless otherwise indicated.

Online supplemental material
Fig. S1 presents ribbon structures of selected parts of BK channels to show the location and size of an arginine sidechain compared with a glycine sidechain for the G375R mutation. Fig. S2 shows the currents recorded from Xenopus oocytes that were used to calculate the relative conductances plotted in Fig. 2. Fig. S3 presents G-V curves for macropatch data obtained in the presence of 300 μM Ca2+ to show that the G375R-induced negative shift also occurs in the presence of 300 μM Ca2+. Fig. S4 presents a statistical test showing that the observed percentages of the five different types of channels expressed when mimicking a heterozygous mutation were not significantly different from the theoretical percentages calculated for an assumption of random assembly of mutant and WT subunits.

Results
G375R subunits left shift BK channel activation
To examine what effect a de novo G375R mutation of the pore-forming subunit of BK channels would have on channel function when coexpressed with WT subunits (Fig. 1), we first compared the whole-cell macroscopic currents flowing through large numbers of WT BK channels expressed following injection of WT cRNA into Xenopus oocytes with those currents expressed following injection of a 1:1 mixture of mutant and WT cRNA to mimic a heterozygous mutation. The injection of a blank without cRNA served as a control. Voltages were stepped from a holding potential of −80 mV to more negative and more positive voltages to reveal the voltage dependent activation of the expressed currents (Fig. 2 A and Fig. S2). Voltage steps to +50 mV were required to appreciably activate WT BK currents generated from the injection of WT cRNA. In contrast, following a 1:1 injection of G375R mutant and WT cRNA, the expressed BK currents were hyperactive, being already active at −140 mV, with depolarization further increasing the response (Fig. 2 B and Fig. S2). The aberrant BK channels from the 1:1 injection that are responsible for these hyperactive currents could be either homotetrameric mutant channels, hybrid channels, or both (Fig. 1).

The negative shift in activation for currents expressed from the 1:1 injection was so pronounced that large numbers of aberrant BK channels were open at voltages near the resting membrane potential of −60 mV. If aberrant currents expressed similarly in neurons, the flux of K+ through opened BK channels

Figure 2. BK channels expressed in oocytes following injection of a 1:1 mixture of G375R mutant and WT cRNA activate at greatly left-shifted negative potentials compared with BK channels expressed following injection of only WT cRNA. (A) Whole-cell currents recorded from oocytes with the two-electrode voltage clamp for the indicated injections of cRNA. The whole cell currents were generated by holding the potential at −80 mV and then jumping to voltages ranging from −140 mV to +60 mV in 10 mV increments (1:1) or to +100 mV (WT and blank). (B) Plots of relative conductance versus the voltage of the steps following injection of a 1:1 mixture of mutant and WT cRNA (red diamonds); injection of WT cRNA alone (black circles); or injection of carrier only (purple triangles). The high conductance at negative potentials would act to drive the membrane potential to about −80 mV, the equilibrium potential of K+. Channel activation is left-shifted in the whole cell recordings in this figure compared to the essentially 0 Ca2+ macro patch recordings in Fig. 3 because the resting free Ca2+ in the oocytes of a few micromolar left shifts BK channel activation. The relative cord conductance was calculated from the plotted steady-state currents in Fig. 52 using the step potential minus the reversal potential for the voltage driving force. Mean ± SEM, n = 4 in each case.
The mean \( mV \) per e-fold change compared with \( 23.2 \pm 1.36 \ mV \) (reduced slope were even greater in the presence of \( 300 \text{ mM} \)). The 1:1 injection also decreased the voltage sensitivity of currents following a 1:1 injection when compared with WT (\( P < 0.0547 \)). The \( G375R \)-induced negative shift in voltage activation and reduced slope were even greater in the presence of \( 300 \text{ mM} \ Ca^{2+} \) (Fig. S3). Thus, the large negative shift in voltage activation observed for whole-cell recordings following a 1:1 injection of mutant and WT cRNA (Figs. 2 and S2) was also observed for macropatch recordings under conditions in which the intracellular solution and \( Ca^{2+} \) were controlled (Figs. 3 and S3).

Whole-cell and macropatch recordings are useful to show the average response of many hundreds to thousands of BK channels, but they provide limited information about the properties of the underlying channels, unless all channels are identical, which is unlikely to be the case for a heterozygous mutation, as shown in Fig. 1. To investigate the channels underlying the negative shifts in BK currents in Figs. 2 and 3 for a 1:1 injection of mutant and WT cRNA, the holding potential was \( -80 \ mV \) to elicit tail currents for \( G/G_{\text{max}} \) measurements. For channels expressed following injection of a 1:1 mixture of \( G375R \) mutant and WT cRNA, the holding potential was \(-160 \ mV \) and voltage pulses were from \(-80 \) to \( +240 \ mV \) with \( 20 \ mV \) steps followed by a step back to \(-80 \ mV \) to elicit tail currents for \( G/G_{\text{max}} \) measurements.

**BK channels of five different potential stoichiometries and six different subunit arrangements could theoretically be expressed for a mutation heterozygous with WT**

WT and mutant subunits arising from a heterozygous mutation could potentially assemble into tetrameric BK channels with five different stoichiometries and six different subunit arrangements, each with potentially different properties (Fig. 1; MacKinnon, 1991; Blaine and Ribera, 1998; Niu and Magleby, 2002; Bergendahl et al., 2019; Backwell and Marsh, 2022). If it is assumed that there is equal production of mutant and WT subunits, that subunits assemble randomly to form tetrameric channels, and that each assembled channel has the same probability of reaching the surface membrane, then the channels expressed for a heterozygous mutation would consist of 6% WT channels, 25% comprised of one mutant and three WT subunits, 38% comprised of two mutant and two WT subunits, 25% comprised of three mutant and one WT subunit, and 6% homotetrameric mutant channels (Fig. 1). On this basis, 94%, of the expressed channels could be pathogenic, comprised of 88% hybrid (heterotetrameric) channels of mixed subunits and 6% homotetrameric mutant channels, with only 6% WT channels (Fig. 1; see unrounded percentages in the figure legend).
Assembled channels
The term assembled channels will be used to refer to those BK channels that are assembled and expressed following injection of a 1:1 mixture of G375R mutant and WT cRNA into Xenopus oocytes to mimic a heterozygous mutation. Assembled channels could theoretically include five types of channels with different stoichiometries and six types if subunit arrangement is considered (Fig. 1). An individual assembled channel could then be any one of the six channel types. It is the combined activity of the six potential types of assembled channels that would determine the BK currents for a heterozygous mutation.

Assembled channels display an abnormally wide range of \( V_h \)
To explore if multiple channel types contribute to the negative shift in \( V_h \) resulting from a G375R BK subunit mutation heterozygous with WT (Figs. 2, 3, S2, and S3), we used the unique ability of the patch clamp technique (Hamill et al., 1981; Niu and Magleby, 2002; Gonzalez-Perez et al., 2015) to isolate and record from individual assembled channels expressed following a 1:1 injection of mutant and WT cRNA into Xenopus oocytes. For each of the 33 individual assembled channels studied, single-channel currents were recorded over a range of voltages (Fig. 4 A) to obtain plots of Po-V (Fig. 4 B, red curves). The Po-V plots for the 33 individual assembled channels had a \( V_h \) that ranged from −152 mV to +151 mV, for a span of 303 mV (Fig. 4 B, red curves). This very wide range in \( V_h \) supports the idea that there are multiple types of assembled channels with different properties arising from different subunit compositions (Fig. 1).

Assembled channels include WT, homotetrameric mutant, and hybrid channels
To identify the types of BK channels expressed following a 1:1 injection of mutant and WT cRNA, the Po-V curves of the 33 individual assembled channels in Fig. 4 B were compared with Po-V curves obtained from individual WT and homotetrameric mutant channels. The WT and homotetrameric mutant channels were obtained by single-channel recording after injection of only WT cRNA or only mutant cRNA, respectively. The WT channels had \( V_h \) values that spanned a narrow range from +140 to +176 mV (Fig. 4 C, black Po-V curves), and the homotetrameric mutant channels had \( V_h \) values that ranged from −272 mV to −38 mV (Fig. 4 C, blue Po-V curves). The homotetrameric mutant channels had a surprisingly wide range of \( V_h \) for channels of the presumed identical composition of four mutant subunits, but all were in a far more negative voltage range than WT channels. The reason for such a wide voltage range in \( V_h \) for the homotetrameric mutant channels is not known, but perhaps channels with four mutant subunits can assume different conformations with markedly different activation properties, depending on mutant side-chain orientation (Fig. S1).

To facilitate the identification of the types of assembled channels, the Po-V curves of the individual 33 assembled channels from Fig. 4 B were overlaid on plots of the \( V_h \) ranges of the Po-V control curves for known WT (gray shading) and homotetrameric mutant (blue shading) channels in Fig. 4 D. One of the 33 assembled channels had a Po-V curve that overlapped with the known WT channel controls (Fig. 4 D), suggesting that this assembled channel was WT with four WT subunits. 4 of the 33 assembled channels had Po-V curves that overlapped with the known homotetrameric mutant channel controls (Fig. 4 D), suggesting that these four assembled channels were homotetrameric mutants comprised of four mutant subunits. The remaining 28 assembled channels had Po-V curves that did not overlap with the known WT or homotetrameric mutant channel controls (Fig. 4 D) but fell in between, suggesting that these 28 assembled channels were all hybrid channels comprised of a mix of mutant and WT subunits (Fig. 1). In this study, assembled channels with \( V_h \) values in the ranges of WT, hybrid, and homotetrameric mutant channels will be referred to as WT, hybrid, and homotetrameric mutant channels, respectively, with the understanding that these classifications are based on \( V_h \) values.

For the sampled group of 33 assembled channels, ~3% (1/33) were consistent with WT, ~85% (28/33) with hybrid, and ~12% (4/33) with homotetrameric mutant (Fig. 4, B–D). Fig. 1 predicts ~6% WT, ~88% hybrid, and ~6% homotetrameric mutant channels. Thus, for the G375R mutation heterozygous with WT, both experimental and theoretical considerations suggest that most (85–88%) of the expressed assembled channels will be hybrid, with much smaller fractions of homotetrameric mutant and WT channels. All hybrid and homotetrameric mutant channels displayed negative shifts in activation compared with WT, with the greatest negative shifts for the homotetrameric mutant channels. Consequently, both theoretical and experimental considerations suggest that 94–97% of the BK channels arising from a G375R mutation heterozygous with WT would display aberrant negative shifts in activation, even though only 50% of the subunits synthesized in a cell would be mutant.

Three functional types of hybrid assembled channels
Analysis in Fig. 4 suggested that assembled channels comprised of 85% hybrid channels, 3% WT, and 12% homotetrameric mutant (Fig. 4). The 85% hybrid channels themselves could consist of three or four different functional types based on both subunit composition and arrangement (Fig. 1). If the functional properties of hybrid channels depend only on subunit stoichiometry, then three functional types would be expected, for one, two, or three mutant subunits replacing an equal number of WT subunits in the heterotetrameric hybrid channels. In addition, if hybrid channels comprising of two mutants and two WT subunits displayed different functional properties for adjacent or diagonal subunit arrangement, then four types of hybrid channels might be expected (Fig. 1). To assess the number of functional types of hybrid channels, a histogram of the \( V_h \) values of the 33 assembled channels from Fig. 4 B was plotted in Fig. 5 A as red bars. Histograms of known WT channel controls (black bars) and homotetrameric mutant channel controls (blue bars) from Fig. 4 C are also plotted. The subunit compositions of the homotetrameric mutant and WT channel controls are known and placed above these two types of channels in Fig. 5 A. As expected from Fig. 4 D, four of the assembled channels had \( V_h \) values that overlapped with those of homotetrameric mutant channels, suggesting that they were homotetrameric mutant channels; one assembled channel overlapped with WT, suggesting that it was a WT channel; and the remaining 28 assembled channels had \( V_h \) values that overlapped with both WT and hybrid channel control curves. However, the number of hybrid channels was not well predicted by the available subunit composition data. The reason for such a large deviation from expected distribution is not clear, but it is possible that further experimental design might improve the theoretical predictions.
values falling in between those of homotetrameric mutant and WT channels, suggesting they were hybrid channels (Fig. 1) whose properties were determined by mixtures of mutant and WT subunits.

The $V_h$ values of the 28 hybrid channels fell into three apparent clusters with mean values of about 1.4, 59, and 96 mV (Fig. 5A). Three clusters of $V_h$ for hybrid channels would be expected from Fig. 1 if each mutant subunit replacing a WT subunit added an increment of a negative shift in $V_h$ independent of subunit arrangement. Accordingly, as a working hypothesis, the different subunit combinations for hybrid channels from Fig. 1 have been placed above the three clusters of hybrid

\[ \text{Figure 4. Identifying the types of assembled BK channels expressed following a 1:1 injection of G375R mutant and WT cRNA. (A) Representative single-channel recordings from three different single BK channels: WT channel (top row) following injection of WT cRNA; an assembled channel following 1:1 injection of G375R mutant and WT cRNA, (second row); and a G375R homotetrameric mutant channel following injection of G375R mutant cRNA (third row). Recordings are shown at three different voltages for each channel type. Depolarization activates each channel type with a markedly different $V_h$ of 140 mV for the WT channel, 6 mV for the assembled channel, and −100 mV for the G375R homotetrameric mutant channel. Arrows indicate closed channel current levels. (B) Plots of $P_o$-V for 33 individual assembled channels following a 1:1 injection of mutant and WT cRNA (red diamonds with Boltzmann fits). (C) Plots of $P_o$-V for 12 single G375R homotetrameric mutant channels following injection of G375R mutant cRNA (blue squares with Boltzmann fits). (D) Identifying the types of assembled channels. Blue and gray areas indicate the range of observed $V_h$ values for G375R homotetrameric mutant channels and WT channels, respectively, from C. One of the assembled channels (rightmost red $P_o$-V curve) overlaps with the WT channels (gray shading), indicating that this assembled channel is likely a WT channel with four WT subunits. Four of the assembled channels (four leftmost red $P_o$-V curves) overlap with the G375R homotetrameric mutant channels (blue shading) suggesting that these four assembled channels are G375R homotetrameric mutant channels assembled from four mutant subunits. The remaining 28 assembled channels would be hybrid channels with mixed subunits (Fig. 1), as their $V_h$ values do not overlap with those of either homotetrameric mutant or WT channels.}
Five BK channel types for a heterozygous mutation

Figure 5. Five different types of functional BK channels are observed following injection of a 1:1 mixture of mutant and WT cRNA into Xenopus oocytes. (A) Histograms of $V_h$ values of single-channels from Fig. 4. WT channels expressed after injection of only WT cRNA are indicated by black bars, and G375R homotetrameric mutant channels expressed after injection of only G375R cRNA are indicated by blue bars. The 33 assembled channels expressed after a 1:1 injection of mutant and WT cRNA are indicated by red bars. The one assembled channel in the cluster of black bars is consistent with channels comprised of four mutant subunits. The 28 assembled channels with a $V_h$ of 151.3 mV, which was significantly different from the 96.4 mV hybrid group, $P < 0.0001$, two-tailed t test. The group of four hybrid channels that fell within the homotetrameric (blue) group had a $V_h$ of $-75.0 \pm 22.0$ mV which was significantly different from the 1.4 mV hybrid group, $P = 0.0019$, two-tailed t test. Hypothesized subunit compositions of the three apparent clusters of hybrid channels are depicted by schematics, where black circles indicate WT subunits and blue circles indicate G375R mutant subunits. Subunit structures for the homotetrameric mutant and WT channels are also indicated. (B) The experimentally observed percentages for each of the five types of 33 assembled channels are listed for comparison to the theoretical percentages from Fig. 1. (C) $V_h$ for the five types of assembled channels is approximated by a linear incremental model where each subunit contributes an increment of $V_h$, $V_h(\text{mix}) = (N_M)(-16.7) + (4 - N_M)(40.5)$, where $N_M$ is the number of G375R mutant subunits per channel, $N_W$ is the number of WT subunits per channel, $V_h(\text{mix})$ is $V_h$, as a function of $N_M$, and $-16.7$ and 40.5 mV are the increments of $V_h$ added by each mutant and WT subunit, respectively. Filled red circles are the experimentally observed mean $\pm$ SEM values of $V_h$ for the five types of assembled channels from A, and the dashed line indicates the predicted values. Data are from 0 intracellular Ca$^{2+}$. Histograms of single-channel kinetic parameters have been used previously to investigate the number of β2 regulatory subunits per BK channel (Wang et al., 2002).

The percentages of expression of the five functional types of assembled channels were then calculated from the number of assembled channels in each cluster in Fig. 5 A and presented in Fig. 5 B. The results are consistent with 3% WT channels, 12% homotetrameric mutant channels, and 85% hybrid channels, where 30% of the hybrid channels had one mutant and three WT subunits, 34% had two mutant and two WT subunits, and 21% had three mutant and one WT subunits. These percentages can be compared with the theoretical values in Fig. 1 calculated for equal production, and a random assembly of subunits where 6% were WT channels, 6% were homotetrameric mutant channels, and 88% were hybrid channels, where 25% of the hybrid channels had one mutant and three WT subunits, 38% had two mutant and two WT channels, and 25% had three mutant and one WT channels. Simulation of 1 million groups of 33 assembled channels, assuming equal production of mutant and WT subunits followed by random assembly into tetrameric channels, indicated that the experimentally observed percentages in Fig. 5 B were not significantly different (Fig. S4) from the theoretical predictions in Fig. 1. A lack of significance does not exclude the possibility that limited amounts of preferential production and assembly of mutant and WT subunits also contributed to the differences between observed and predicted percentages, in addition to the large variability arising from the random assembly of subunits that are characterized in Fig. S4.

Whereas the observation of three apparent clusters of $V_h$ values for the hybrid assembled channels (Fig. 5) is consistent
with theoretical predictions (Fig. 1), peaks can occur by chance alone in histograms of binned data of limited sample size (Miller et al., 1978). Consequently, additional experiments would be needed to determine whether $V_h$ values for hybrid channels can consistently be resolved into three peaks, but an observation of such distinct peaks is not required for support of Fig. 1, as variability in $V_h$ among channels of the same type (McManus and Magleby, 1991) might be sufficient to obscure distinct peaks.

**Linear incremental model for the contributions of mutant and WT subunits to $V_h$**

WT BK channels comprised of four WT subunits had a mean $V_h$ of about 160 mV, and homotetrameric mutant BK channels comprised of four mutant subunits had a mean $V_h$ of about −80 mV (Fig. 4 and Fig. 5A). As all four subunits contribute to the gating of BK channels (Rothberg and Magleby, 2000; Horrigan and Aldrich, 2002; Niu and Magleby, 2002), these observations suggest that each WT subunit contributes an increment of a positive shift to $V_h$, and that each mutant subunit contributes an increment of negative shift. In support of this hypothesis, a linear incremental model in which each WT subunit in a channel added +40.5 mV to $V_h$, and each mutant subunit added −16.7 mV provided an approximate description of the observed $V_h$’s for the five types of assembled channels (Fig. 5C dashed line; model in the figure legend). The net effect of replacing a WT subunit with a mutant subunit was a −57.2 mV shift in $V_h$ to account for the loss of the positive shift contributed by the removed WT subunit and the addition of the negative shift contributed by the added mutant subunit.

The mechanism by which each subunit with a G375R mutation provides a net −57.2 mV shift in $V_h$ is not known, but is unlikely to involve the addition of positive charge by arginine because the G375D mutation (G310D in Chen et al., 2014) which adds negative charge also gives a negative shift in $V_h$, but of smaller magnitude. That the left shift in $V_h$ increases with side chain volume, where arginine > aspartate > glycine suggests that larger side chain volumes at the hinge position of G375 may progressively decrease the energy barrier for opening.

**Figure 6.** Each replacement of a WT subunit with a G375R mutant subunit in a BK channel acts to both left shift $V_h$ to more negative potentials and decrease single channel conductance. (A) Single-channel currents were recorded from single channels at +100 mV after injecting the indicated cRNA. (B) Single-channel conductance $g$ at +100 mV decreases as the number of mutant subunits increases. Mean ± SEM single-channel conductance for WT channels was $312 ± 6$ pS (n = 7), decreasing to $245 ± 6$ pS for hybrid channels ($n = 25$, $P < 0.0001$), and further decreasing to $190 ± 14$ pS for G375R homotetrameric mutant channels ($n = 8$, $P = 0.0003$). Mean ± SEM WT and G375R homotetrameric mutant channels were those channels expressed after injecting only WT cRNA or only G375R mutant cRNA, respectively. Hybrid channels were identified as in Fig. 4D. (C) Plot of single channel conductance at +100 mV vs. $V_h$ for BK channels expressed following injection of the indicated cRNA. The linear regression line plots single channel conductance $g$ vs. $V_h$, where $g = g(0) + sV_h$, where $g(0) = 231.9 ± 3.9$ pA, and $s = 0.408 ± 0.0375$ pS/mV ($P < 0.0001$ for slope significantly different from 0, $R = 0.86$). (D) Single-channel conductance for the five types of assembled channels is approximated by a linear incremental model, where each subunit contributes an increment of conductance, $g_{\text{increment}} = (N_M)(46.1 pS + 4 - N_M)(76.5 pS)$, where $N_M$ is the number of G375R mutant subunits per channel, $4 - N_M$ is the number of WT subunits per channel, $g(0)$ is single channel conductance as a function of $N_M$, and 46.1 pS and 76.5 pS are the increments of $g$ added by each mutant and WT subunit, respectively. The red dashed line indicates the predicted values of $g$. The homotetramer mutant (blue) and WT (black) conductance bars are from B. The three red hybrid conductance bars plot the mean ± SEM of $g$ for the three groups of hybrid channels identified in Fig. 5A, where $g$ measurements were available at +100 mV, giving: 209.0 ± 7.5 pS, $n = 7, 246.0 ± 7.7$ pS, $n = 8$, and 269 ± 2.6 pS, $n = 10$. For the four possible pairs of adjacent single-channel conductance bars in D, P values were calculated for $g$ of the right conductance bar in each pair being significantly different than $g$ for the left, and were 0.26, 0.0034, 0.0064, 0.0002, respectively; two-tailed t tests.
Dual action of G375R subunits on $V_h$ and single channel conductance

In addition to the negative shift in activation induced by replacing WT subunits with G375R mutant subunits (Figs. 2, 3, 4, and 5), replacing WT subunits with mutant subunits also decreased single-channel conductance (Fig. 6, A and B). The mean single-channel conductance of WT channels was 312 ± 4 pS. This decreased to 245 ± 6 pS for hybrid channels and further decreased to 190 ± 14 pS for homotetrameric mutant channels. These decreases were significant (Fig. 6 legend). Hence, single-channel conductance decreased as mutant subunits replaced WT subunits. The decreased single-channel conductance may arise from the larger volume and positive charge of the mutant arginine side chains replacing the single hydrogen atom of the glycine side chains on one or more of the S6 segments lining the conductance pathway of the BK channel (Fig. S1). The added volume of the side chains could decrease the volume of the inner vestibule and the added positive charge may act to repel K$^+$ from the inner cavity. Both actions can reduce single-channel conductance in BK channels (Brelidze et al., 2003; Geng et al., 2011; Nimigean et al., 2003).

To examine the relationship between single-channel conductance and $V_h$, the single-channel conductance for each channel was plotted against $V_h$ for the same channel in Fig. 6 C for the indicated channel types. A linear relationship was observed (Fig. 6 C; $R = 0.86$, $P < 0.0001$). When taken together, the data in Fig. 6, B and C, are consistent with the idea that replacing a WT subunit with a mutant subunit adds both an increment of negative voltage shift to $V_h$ and a step decrease in single-channel conductance. Thus, at the single-channel level, the heterozygous G375R mutation acts simultaneously as a GOF mutation to shift voltage activation to more negative voltages (Figs. 4 and 5) and as a DOF mutation to decrease single-channel conductance (Fig. 6). At the whole-cell and macropatch level, the increase in currents from the GOF negative shift in activation would dominate the DOF reduction in single-channel conductance, producing large left-shifted currents (Figs. 2, 3, S2, and S3). The DOF in conductance would act to decrease the consequences of the negative shift in activation.

Linear incremental model for the contributions of mutant and WT subunits to single-channel conductance

A linear incremental model in which each WT subunit contributes 76.5 pS to single channel conductance and each mutant subunit contributes 46.1 pS could approximate the single-channel conductance for the five types of assembled channels (Fig. 6 D dashed line; model in the figure legend).

WT, hybrid, and G375R homotetrameric mutant BK channels are expressed in the HEK293 cell expression system

Liang et al. (2019) reported that no potassium currents were recorded from excised macropatches following transfection of HEK293T cells with plasmids coding for G375R mutant BK subunits. In contrast, we found that functional single homotetrameric mutant channels were readily expressed following injection of Xenopus oocytes with cRNA coding for mutant BK G375R subunits (Fig. 4, A, and C). To examine if these differences in channel expression were due to differences in expression systems, we transfected HEK293 cells with WT cDNA to generate WT channels, with mutant G375R cDNA to generate homotetrameric mutant channels, and with a 1:1 mixture of WT and mutant cDNA to mimic the generation of channels that would normally arise from a heterozygous mutation. We found that the transfected HEK293 cells displayed robust whole-cell BK currents for each of the three different types of transfections (Fig. 7 A). The macroscopic G-V curves from HEK293 cells for transfection with WT cDNA alone or 1:1 transfection with WT and mutant cDNA were very similar to the macroscopic G-V curves we obtained using the Xenopus oocyte expression system (compare Fig. 7 with Fig. 3). We also found that the whole-cell current from G375R homotetrameric mutant channels expressed in HEK293 cells had a $V_h$ value within the broad range of $V_h$ values of single-channel recordings from patches of membrane excised from Xenopus oocytes injected with G375R mutant mRNA (Fig. 4).

Hence, in both Xenopus oocytes and HEK293 expression systems, we observed functional G375R homotetrameric mutant currents that displayed a marked negative shift in activation consistent with a GOF mutation. This contrasts with the observations of Liang et al. (2019), who did not observe BK channel currents for G375R transfection of HEK293T cells. They reported G375R as a LOF mutation. The reason for the difference in observations is not known, but we found that the viability of Xenopus oocytes injected with G375R cRNA was reduced (see Materials and methods).

Whole-cell and macropatch currents can obscure mechanism

The narrow error bars for the macro currents recorded from whole oocytes and excised macropatches (Figs. 2, 3, S2, and S3) following injection of WT cRNA or a 1:1 injection of mutant and WT cRNA, indicated that the mean responses recorded from hundreds to thousands of channels were repeatable. Narrow error bars for mean responses do not necessarily indicate that the individual channels underlying the mean response have near identical properties, as this was not the case for the macro currents following 1:1 injection of mutant and WT cRNA, where individual assembled channels following the 1:1 injection had a very wide range of $V_h$ that spanned over 300 mV (Fig. 4 B). In spite of this wide range for individual channels, the mean G-V data following the 1:1 injections were reasonably well described by a single Boltzmann function (Fig. 3 B, blue line through red diamonds), but were better described (red line) by assuming five underlying channel types, as described in the legend to Fig. 3.

Discussion

The heterozygous G375R BK channel variant has been associated with a devastating human phenotype that includes malformation syndrome and severe neurological and developmental disorders (Liang et al., 2019). This variant has only appeared in the human population when heterozygous with WT, perhaps because a homozygous G375R genotype may not permit viability because of the extreme GOF phenotype we observed for homotetrameric mutant channels. To gain insight into the
pathogenicity of this variant, currents from whole cells, macropatches, and single channels were recorded from BK channels expressed following a 1:1 injection of G375R mutant and WT cRNA to mimic a G375R mutation heterozygous with WT.

Recordings from whole cells and macropatches, both of which contain many hundreds to thousands of channels, indicated that the V$_h$ of the current following a 1:1 injection, was left shifted to more negative potentials by about −120 mV compared with WT currents (Figs. 2, 3, S2, and S3). The aberrant BK channels underlying the negative shifts in activation would lead to a much greater fraction of BK channels being open at negative membrane potentials, including at potentials near the resting potential (Figs. 2, 3, S2, and S3), which would oppose cellular depolarization, altering cellular function.

To explore the underlying molecular basis for the aberrant current activation associated with the heterozygous G375R variant, we used high-resolution single-channel recording to isolate and characterize the BK channels that are assembled and expressed following injection of a 1:1 mixture of G375R mutant and WT cRNA. Theoretical considerations based on equal production and a random assembly of subunits suggest there could be multiple types of assembled channels with different subunit combinations (MacKinnon, 1991; Blaine and Ribera, 1998; Bergendahl et al., 2019; Backwell and Marsh, 2022; Fig. 1). Consistent with this possibility, our analysis suggested that five different types of functional BK channels were expressed: 3% were consistent with WT, 12% with homotetrameric mutant, and 85% with three different types of hybrid channels of mixed subunits (Fig. 4; and Fig. 5, A and B). The percentages of expression of these five types of functionally assembled channels were not significantly different (Fig. S4) from the theoretical predictions of Fig. 1 based on equal production and a random assembly of subunits. This suggests, within the limits of experimental variability (Fig. S4), that the processes involved in subunit production, assembly, and expression do not distinguish significantly between mutant and WT alleles and subunits, so the phenotypic differences arise at the functional level of individual channels.

The three types of hybrid channels comprising 85% of the expressed assembled channels had properties falling between those of mutant and WT channels that varied with their apparent subunit composition (Figs. 4, 5, and 6). 97% of the assembled channels, all except for the 3% WT, displayed both GOF negative shifts in activation and smaller DOF reductions in single-channel conductance (Figs. 4, 5, and 6). Hence, most of the channels expressed for the heterozygous G375R mutation displayed aberrant properties. The values of V$_h$ and single channel conductance for each of the five types of functional assembled channels could be predicted with a linear incremental model in which each mutant and WT subunit in each of the five types of functional assembled channels acted relatively independently to contribute increments of both V$_h$ and single-channel conductance to the molecular phenotype of the channel (Fig. 5 C and Fig. 6 D; the models are in the figure legends).
A potential mechanism to produce the five functional types of assembled channels for a heterozygous G375R BK channel mutation, then, is equal production and a random assembly of mutant and WT subunits into channels of five different subunit compositions, where the \( V_h \) and single-channel conductance of each channel type are determined by independent contributions from each of the four subunits in a channel. Each WT subunit adds increments of 40.5 mV to \( V_h \) and 76.5 pS to single-channel conductance, and each mutant subunit adds increments of −16.7 mV to \( V_h \) and 46.1 pS to single-channel conductance (Fig. 5 C and Fig. 6). Whereas a linear incremental model could provide reasonable descriptions of the data, further study is likely to reveal added complexity.

We were surprised to find that the mean macroscopic G-V curve following a 1:1 injection of G375R mutant and WT cRNA could also be well described by a single Boltzmann function (Fig. 3) as the G-V curve arose from the sum of currents from five different types of BK channels with markedly different properties (Figs. 4, 5, and 6). Consequently, an observation that a macroscopic G-V curve is well described by a single Boltzmann function does not necessarily exclude the possibility that the macroscopic G-V curve arises instead from multiple types of channels with different properties. The description with a single Boltzmann function may be possible in this case because the percentages of the five types of contributing channels first increase and then decrease (Fig. 1 and Fig. 5 B), helping to fill in and smooth the G-V curve. Paradoxically, the single-Boltzmann shape of the G-V curves for 1:1 injections and transfections (Fig. 3 and Fig. 7) provides additional evidence for the assembly and expression of hybrid channels, as there would be two clearly separable Boltzmann components in the 1:1 G-V curves arising from homotetrameric mutant and WT channels if the G375R and WT subunits did not coassemble to form three types of hybrid channels with intermediate properties to fill in and smooth the G-V curve.

For classification with regard to genetic disease (Backwell and Marsh, 2022), we suggest that the heterozygous G375R mutation be labeled as an assembly-mediated dominant GOF mutation: assembly-mediated to indicate that mutant and WT subunits assemble into multiple types of functional tetrameric BK channels, dominant because assembled channels with one or more mutant subunits display pathogenic properties, and GOF because less depolarization is required to activate BK channels with one or more mutant subunits. The net result is that 94–97% of the expressed channels display aberrant activation (Figs. 1 and 5). A dominant GOF phenotype has been described previously for the heterozygous G88R mutation of the TASK-4 K⁺ channel in the heart, based on a study using whole-cell currents (Friedrich et al., 2014).

An assembly-mediated dominant-GOF mutation for heterozygous G375R can be compared to the well-known dominant-negative effect observed for some types of protein complexes and channelopathies (Fink et al., 1996; Ribera et al., 1996; Silberberg et al., 2005; Reed et al., 2016; Du et al., 2020; Hichri et al., 2020; Backwell and Marsh, 2022). Both can cause disease through a disproportionate fraction of the channels affected. In the first case, there is a GOF in channels with one or more mutant subunits, and in the second there is a complete or partial loss of function of channels associated with, typically, one or more mutant subunits.

Whereas an assembly-mediated dominant GOF classification of the heterozygous G375R mutation is useful to suggest the potential underlying basis of genetic disease, as it describes the net functional cellular phenotype, the molecular phenotype is more complex. The action of the heterozygous G375R mutation is to generate five types of channels, four of which simultaneously display a GOF in activation and a smaller DOF in single-channel conductance. The GOF dominates the response at the level of cellular currents.

The random assembly of mutant and WT subunits into tetrameric BK channels when mimicking a heterozygous mutation led to multiple types of genetic dominance at the level of the molecular phenotype when viewed with single-channel recording. Codominance was observed for the WT and homotetrameric mutant channels, and partial dominance was observed for the three types of hybrid channels. Support for codominance was that the homotetrameric mutant and WT channels expressed following injection of only mutant or WT cRNA (Figs. 4, 5, and 6). Support for partial dominance was that the three types of expressed hybrid channels had individual values of \( V_h \) and single channel conductance that were distinct from each other and fell in the range between those of homotetrameric mutant and WT channels (Fig. 5 C and Fig. 6 D). The levels of partial dominance in the linear incremental model (Fig. 5 C and Fig. 6 D) were determined by the numbers of mutant and WT subunits per channel acting independently of one another, rather than by mutant subunits altering the function of WT subunits, as in some classical descriptions of genetic dominance. It is remarkable that a single base pair substitution in one allele of a pair of alleles that encode for the \( \alpha \) subunit of BK channels results in the expression of WT BK channels plus four aberrant types of BK channels, each with different molecular phenotypes that simultaneously display a dominant GOF in \( V_h \) and a lessor DOF in single-channel conductance, with the five types of functional channels displaying either codominance or one of three levels of partial dominance for both \( V_h \) and single channel conductance.

How do our observations of independent interactions of the G375R mutant and WT pore-forming subunits compare to interactions of these subunits with regulatory subunits? Wang et al. (2002) found that single BK channels comprised of four subunits could be associated with zero to four regulatory \( \beta_2 \)-subunits per channel, with each \( \beta_2 \)-subunit giving incremental changes in \( V_h \). Hence, both mutated and WT pore-forming subunits (Fig. 5 C; Niu and Magleby, 2002) and the non-pore-forming regulatory \( \beta_2 \)-subunits (Wang et al., 2002) can give incremental changes in gating, but such incremental changes per subunit are not necessarily universal, so each type of subunit will need to be assessed. A single BK channel comprised of four pore-forming subunits can also include up to four regulatory \( \gamma_1 \)-subunits, but a single \( \gamma_1 \)-subunit per channel is sufficient to induce the full gating shift induced by \( \gamma_1 \)-subunits (Gonzalez-Perez et al., 2014, 2018).

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Five BK channel types for a heterozygous mutation

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Devising therapies for the de novo G375R heterozygous mutation will be challenging. Theoretical and experimental observations (Figs. 1, 4, and 5) suggest that most (94–97%) of the channels for a mutation heterozygous with WT would be pathogenic, with only 3–6% of the channels WT. The pathogenic channels would consist of multiple channel types, each with large differences in activation properties and smaller differences in conductance (Figs. 4, 5, and 6). Therapies to block or inactivate the pathogenic channels would ideally silence the multiple types of pathogenic channels while leaving any WT channels intact. Even if such selective blockers could be devised, it is unlikely that the remaining 3–6% of WT channels would be sufficient to restore normal cellular function. Effective therapies will likely require replacing or silencing the mutant alleles or preventing mutant subunits from assembling with themselves and WT subunits if/when such techniques become practical in humans.

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Figure S1. Cryo-EM ribbon structures of the Ca$^{2+}$-free human BK channel (PDB ID: 6V3G; Tao and MacKinnon, 2019) showing the location of the mutant arginine side chain G375R. (A) BK channel structure with the front and back pore-forming α subunits removed. The arginine side chain of the G375R mutation is shown in space-filling format on the back side of S6 away from the conduction pathway on the red-colored subunit. Carbon atoms are gray, nitrogen atoms are blue, and hydrogen is not indicated. (B) Structure after rotating 180° so that the location (blue mark) of the WT glycine side chain of G375 is visible on the back side of S6 on the yellow subunit. (C) Six possible orientations of the mutant arginine side chain are shown, generated by ChimeraX software (www.cgl.ucsf.edu/chimerax). The mutation G375D also produces a left shift in activation (G310D in Chen et al., 2014), but of half the magnitude of G375R (Fig. 3). This raises the possibility that the left shift may be related to the bulk of the substituted side chain rather than the specific charge, which is reversed for these two mutations.
Figure S2. BK currents in Xenopus oocytes following injection of a 1:1 mixture of G375R mutant and WT cRNA activate at greatly left-shifted negative voltages compared with BK currents following injection of only WT cRNA. I-V plots for the indicated injections of cRNA and blank. Two-electrode whole-cell voltage clamp. These currents were used to calculate the relative conductance plotted in Fig. 2. Protocol details are in Fig. 2 legend. The amount of WT cRNA injected was 25 times greater than for the injection of the 1:1 mixture of G375R mutant and WT cRNA so that the initial deviation of WT currents from the baseline could be readily detected. Mean ± SEM, n = 4 in each case.
The aberrant negative shift in $V_h$ for BK currents expressed following a 1:1 injection of G375R mutant and WT cRNA is also observed with 300 $\mu$M intracellular Ca$^{2+}$. (A) Currents recorded from inside-out macro patches of the membrane with 300 $\mu$M Ca$^{2+}$ at the inner membrane surface. For WT channels, voltage pulses were from $-160$ to 180 mV with 20 mV steps from a holding potential of $-160$ mV. Conductance was measured from tail currents after stepping to $-160$ mV. For channels expressed following injection of a 1:1 mixture of G375R mutant and WT cRNA, voltage pulses were from $-200$ to 180 mV with 20 mV steps from a holding potential of $-200$ mV. Conductance was measured from tail currents after stepping to $-200$ mV. The dashed line indicates the level of 0 current. In the presence of 300 $\mu$M intracellular Ca$^{2+}$, more than half of the channels expressed following the 1:1 injection of G375R mutant and WT cRNA remain open at $-200$ mV. The macrocurrents are the average response from many hundreds of BK channels in each macro patch. (B) G-V plots with 300 $\mu$M Ca$^{2+}$ following injection of WT cRNA alone (black circles) or a 1:1 injection of mutant and WT cRNA (red diamonds). The mean $V_h$ for WT currents was $-7.2 \pm 3.3$ mV, $n = 8$, shifting to $-205 \pm 9.6$ mV for BK currents expressed from a 1:1 mixture of mutant and WT cRNA, producing a left shift of $-198$ mV with 300 $\mu$M Ca$^{2+}$ at the inner membrane surface, $n = 6$. This can be compared with a left shift of $-120$ mV with 0 Ca$^{2+}$ (Fig. 3). Hence, the aberrant left shift in $V_h$ induced by a 1:1 injection of mutant and WT cRNA occurs in the presence and absence of intracellular Ca$^{2+}$.
The experimentally observed percentages of the five types of assembled channels expressed following a 1:1 injection of G375R mutant and WT cRNA did not differ significantly from the theoretical percentages calculated assuming random assembly of G375R mutant and WT subunits. Simulation was used to determine if the percentages of observed and predicted channel types (Fig. 5 B and Fig. 1) were significantly different by examining whether the observed numbers of channels of each type in the experimental group of 33 assembled channels (Fig. 5 A) were significantly different from the expected numbers based on random assembly of subunits (this figure), as it is the numbers that determine the percentages. The expected numbers could not be calculated directly because of the random assembly of subunits but could be specified in terms of simulated discrete probability distributions, one for each of the five channel types (see Materials and methods). This figure plots the five discrete probability distributions, where the specific channel type (stoichiometry) for each distribution is indicated by a schematic, where black circles are WT subunits and blue circles are mutant subunits. Each one of the five discrete probability distributions plots the probability of observing 0, 1, 2, 3...33 assembled channels of the type indicated for that distribution in a group of 33 channels randomly assembled from mutant and WT subunits. The five discrete probability distributions were generated by first simulating 10^6 groups of 33 randomly assembled channels (see Materials and methods and Fig. 1) and then binning the number of each type of assembled channel in each group of 33 channels into five frequency histograms, one for each channel type. The bins in each frequency histogram were then divided by 10^6, the total counts in each frequency histogram, to normalize each frequency histogram to discrete probability distributions with an area of 1.0. To illustrate with some examples, the topmost discrete probability distribution, which gives the probabilities of observing different numbers of WT channels in a group of 33 assembled channels, indicates that the probability of observing 0 WT channels in a group of 33 randomly assembled channels is 0.119. The probability of observing 1 WT channel is 0.262, and the probability of observing 2 WT channels is 0.278, etc. Thus, 11.9, 26.2, and 27.8% of the groups of 33 assembled channels would have 0, 1, or 2 WT channels, respectively. The arrow at 2.06 WT channels indicates the mean number of WT channels per group of 33 assembled channels from the analysis of the 10^6 groups. The expected mean number of WT channels per group of 33 channels can also be calculated directly from the percentages in Fig. 1, where 6.25% of the 33 assembled channels on average would be WT, giving a mean of 2.06 WT assembled channels per group of 33 assembled channels from the analysis of the 10^6 groups. The expected mean number of WT channels per group of 33 assembled channels can also be calculated directly from the percentages in Fig. 1, where 6.25% of the 33 assembled channels on average would be WT, giving a mean of 2.06. The discrete probability distributions for each of the four other types of assembled channels are also presented and can be interpreted in a similar manner. Our experimental observations of the number of assembled channels of each type in the experimental group of 33 assembled channels (Fig. 4 D, and Fig. 5, A and B) are indicated by the locations of the red histogram bars on each abscissa of the discrete probability distributions. The amplitude of the red bars gives the probability of observing the indicated number of channels of that type assuming a random assembly of subunits. For example, the probability for our observation of 1 WT channel in the experimental group of 33 assembled channels was 0.262, and the probabilities for our observations of 10, 11, 7, and 4 assembled channels for the other indicated types of assembled channels are also presented and can be interpreted in a similar manner. Our experimental observations of the number of assembled channels of each type in the experimental group of 33 assembled channels (Fig. 4 D, and Fig. 5, A and B) are indicated by the locations of the red histogram bars on each abscissa of the discrete probability distributions. The amplitude of the red bars gives the probability of observing the indicated number of channels of that type assuming a random assembly of subunits. For example, the probability for our observation of 1 WT channel in the experimental group of 33 assembled channels was 0.262, and the probabilities for our observations of 10, 11, 7, and 4 assembled channels for the other indicated types of assembled channels in the experimental group of 33 assembled channels were 0.118, 0.128, 0.147, and 0.0961, respectively. All of these probabilities for the observed numbers of assembled channels of each type were >0.05, and none of the observed numbers of assembled channel types fell within the 0.05 summed cumulative probability of the left and right tails of the discrete probability distributions, indicating that the experimental observations for the number of the five types of assembled channels were not significantly different (Colquhoun, 1971) from the theoretical predictions based on an assumption of random subunit assembly. A lack of significant difference does not exclude that there may be underlying differences between experimental and theoretical predictions, which could be revealed by larger sample sizes.